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(54) **Pentafluorophenylazolopyrimidines**

Pentafluorophenylazolopyrimidine

Pentafluorophénylazolopyrimidines

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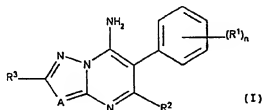
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Description

BACKGROUND OF THE INVENTION

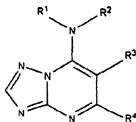
[0001] This invention relates to certain triazolopyrimidine compounds, a process for their preparation, compositions containing such compounds, a method for combating a fungus at a locus comprising treating the locus with such compounds and their use as fungicides.

[0002] EP-A-0071792 discloses compounds of formula I



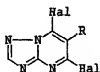
in which R¹ represents alkyl, halogen, alkoxy, cyano, cycloalkyl, aryl, aryloxy, arylthio, aralkyl, arylthio, arylalkyl, arylalkyloxy or arylalkylthio each optionally substituted by halogen or alkoxy; or (R¹)_n represents a benzene, indane or tetrahydronaphthalene ring fused with the phenyl ring, aromatic moieties in the above groups being optionally substituted by alkyl, alkoxy, halogen or cyano; n is 1 or 2; R² and R³ are each hydrogen, alkyl or aryl, A represents a nitrogen atom or a CR⁴ group, and R⁴ is as R² but can also be halogen, cyano or alkoxycarbonyl or together with R³ can form an alkylene chain containing up to two double bonds. The compounds are said to be active against various phytopathogenic fungi, especially those of the phycomycete class. However evidence of fungicidal activity is only provided for these compounds against *Plasmopara viticola*, a member of the oomycete class of fungi.

[0003] EP 550113-A2 discloses compounds of the general formula



in which R¹ represents an optionally substituted alkyl, alkenyl, alkadienyl, cycloalkyl, bicycloalkyl or heterocyclyl group; R² represents a hydrogen atom or an alkyl group; or R¹ and R² together with the interjacent nitrogen atom represent an optionally substituted heterocyclic ring; R³ represents an optionally substituted aryl group; and R⁴ represents a hydrogen or halogen atom or a group -NR⁵R⁶ where R⁵ represents a hydrogen atom or an amino, alkyl, cycloalkyl or bicycloalkyl group and R⁶ represents a hydrogen atom or an alkyl group. These compounds are said to be active against fungi which are members of the ascomycetes class such as *Venturia inaequalis* and of the hyphomycetes class such as *Alternaria solani* and *Botrytis cinerea*.

[0004] WO-A 94/20501 discloses 5,7-dihalogenated triazolopyrimidines of the general formula

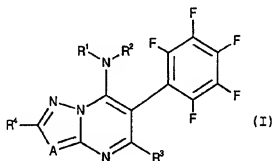


in which R represents an optionally substituted aryl group. These compounds are said to be active against fungi.

[0005] Intermediate compound ethyl pentafluorophenyl malonate is known from J. Org. Chem. USSR, 6, 761 (1970) and GB-A 901 880.

Summary of the invention

[0006] The present invention provides novel pentafluorophenyl substituted azolopyrimidines of formula I



In which

- R¹ and R² each independently represent hydrogen or C₁-C₁₀-alkyl, C₂-C₁₀-alkenyl, C₂-C₁₀-alkynyl, C₄-C₁₀-alkadienyl, C₆-, C₁₀- or C₁₄-aryl, heteroaryl having 5 or 6 ring atoms selected from carbon, nitrogen, oxygen and sulphur, at least one of which being nitrogen, oxygen or sulphur, C₃-C₈-cycloalkyl, saturated heterocyclyl having 5 or 6 ring atoms selected from carbon, nitrogen, oxygen and sulphur, at least one of which being nitrogen, oxygen or sulphur, or
- R¹ and R² together with the adjacent nitrogen atom represent a heterocyclic ring, wherein each group is optionally substituted by one or more halogen atoms or nitro, cyano, C₁₋₆-alkyl, C₃₋₆-cycloalkyl, C₃₋₆-cycloalkenyl, C₁₋₆-haloalkyl, C₃₋₆-halocycloalkyl, C₁₋₆-alkoxy, C₁₋₆-haloalkoxy, phenyl, halo- or di-halo-phenyl or pyridyl.
- R³ represents a halogen atom or a group -NR⁵R⁶, wherein R⁵ and R⁶ each independently have one of the meanings given for R¹ and R²,
- R⁴ represents hydrogen or C₁-C₁₀-alkyl or C₆-, C₁₀- or C₁₄-aryl, and
- A represents N or CR⁷, wherein R⁷ has the meaning given for R⁴.

[0007] It is another object of the invention to provide a new process for the preparation of these novel compounds and to provide novel intermediates which are prepared during this new process.

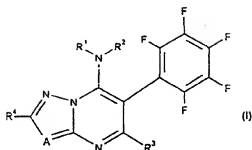
[0008] It is another object of the invention to provide a fungicidal composition which comprises a carrier, and as active agent, at least one compound of formula I according to the invention.

[0009] It is another object of the invention to provide a method of combating fungus at a locus which comprises treating the locus with a compound of formula I according to the invention or with a composition comprising a compound of formula I according to the invention.

[0010] These and other objects and features of the invention will become more apparent from the detailed description set forth hereinbelow.

DETAILED DESCRIPTION OF THE INVENTION

[0011] It has surprisingly been found out that the novel pentafluorophenylazolopyrimidines of formula I



in which A and R¹ through R⁴ have the meaning given above for formula I unexpectedly show excellent fungicidal activity against a broad range of fungi and are obtainable through a process which includes four synthetic steps from commercially accessible educts.

[0012] In general terms, unless otherwise stated, as used herein the term halogen atom may denote a bromine, iodine, chlorine or fluorine atom, and is especially a bromine, chlorine or fluorine atom.

[0013] Optionally substituted moieties may be unsubstituted or have from one up to the maximal possible number of substituents. Typically, 0 to 2 substituents are present.

[0014] In general terms, unless otherwise stated herein, the terms alkyl, alkenyl, alkynyl, alkyldienyl as used herein with respect to a radical or moiety refer to a straight or branched chain radical or moiety. As a rule, such radicals have up to 10, in particular up to 6 carbon atoms. Suitably an alkyl moiety has from 1 to 6 carbon atoms, preferably from 1 to 3 carbon atoms. A preferred alkyl moiety is an ethyl or especially a methyl group. Suitably an alkenyl moiety has from 2 to 6 carbon atoms. A preferred alkenyl moiety is allyl or especially a 2-methylallyl group.

[0015] In general terms, unless otherwise stated herein, the term aryl, as used herein with respect to a radical or moiety refers to an aryl group having 6, 10 or 14 carbon atoms, preferably 6 or 10 carbon atoms. In particular phenyl being optionally substituted by one or more halogen atoms, nitro, cyano, alkyl, preferably C₁₋₆ alkyl, alkoxy, preferably C₁₋₆ alkoxy.

[0016] In general terms, unless otherwise stated herein, the term heteroaryl, as used herein with respect to a radical or moiety refers to a heteroaryl group having 5 or 6 ring atoms selected from carbon, nitrogen, oxygen and sulphur, at least one of which being nitrogen, oxygen or sulphur.

[0017] In general terms, unless otherwise stated herein, the term cycloalkyl, as used herein with respect to a radical or moiety refers to a cycloalkyl group having 3 to 8 carbon atoms, preferably 5 to 7 carbon atoms, in particular cyclohexyl being optionally substituted by one or more halogen atoms, nitro, cyano, alkyl, preferably C₁₋₆ alkyl, alkoxy, preferably C₁₋₆ alkoxy.

[0018] In general terms, unless otherwise stated herein, the term heterocyclyl, as used herein with respect to a radical or moiety refers to a saturated heterocyclyl group having 5 or 6 ring atoms selected from carbon, nitrogen, oxygen and sulphur, at least one of which being nitrogen, oxygen or sulphur being optionally substituted by one or more halogen atoms, nitro, cyano, alkyl, preferably C₁₋₆ alkyl, alkoxy, preferably C₁₋₆ alkoxy, in particular pyrrolidinyl, piperazinyl, piperidinyl, piperazinyl or morpholin-4-yl.

[0019] The invention especially relates to compounds of the general formula I in which any alkyl part of any of the groups R¹, R², R³ and R⁴, which may be straight chained or branched, contains up to 10 carbon atoms, preferably up to 6 carbon atoms, more preferably up to 6 carbon atoms, any alkenyl or alkynyl part of any of the substituents R¹ to R⁶ contains up to 10 carbon atoms, preferably up to 9 carbon atoms, more preferably up to 6 carbon atoms, any cycloalkyl part of any of the substituents R¹ to R⁶ contains from 3 to 10 carbon atoms, preferably from 3 to 8 carbon atoms, more preferably from 3 to 6 carbon atoms, any heterocyclic ring formed from R¹ to R⁶ with the adjacent nitrogen atom, contains from 3 to 10 ring members, preferably from 3 to 6 carbon atoms, and any aryl part of any of the substituents R¹ to R⁶ contains 6, 10 or 14 carbon atoms, preferably 6 or 10 carbon atoms, and in which each optionally substituted group independently is substituted by one or more halogen atoms or nitro, cyano, alkyl, preferably C₁₋₆ alkyl, cycloalkyl, preferably C₃₋₆ cycloalkyl, cycloalkenyl, preferably C₃₋₆ cycloalkenyl, haloalkyl, preferably C₁₋₆ haloalkyl, haloalkoxyalkyl, preferably C₃₋₆ haloalkoxyalkyl, alkoxy, preferably C₁₋₆ alkoxy, haloalkoxy, preferably C₁₋₆ haloalkoxy, phenyl, halo- or dihalo-phenyl or pyridyl groups. Any alkyl, alkenyl or alkynyl group may be linear or branched. A 4- to 6-membered heterocyclic group may be any heterocyclic group with 4 to 6 ring atoms, interrupted by one or more heteroatoms selected from sulfur, nitrogen, and oxygen, preferably oxygen. A halogen atom suitably denotes a fluorine, chlorine or bromine atom.

[0020] The invention especially relates to compounds of formula I in which R¹ represents a C₁₋₁₀ alkyl, C₃₋₈ cycloalkyl, C₃₋₈ cycloalkyl, C₁₋₆ alkyl, C₁₋₁₀ alkoxy, C₁₋₆ alkyl or phenyl group.

[0021] Preferably R¹ represents a C₁₋₈, suitably C₁₋₆, alkyl group, especially a branched alkyl group, more especially

secondary and tertiary alkyl groups as secondary butyl, tertiary butyl and tertiary amyl groups.

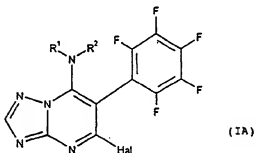
[0022] The invention further especially relates to compounds of formula I in which R¹ and R² each independently represent a hydrogen atom, a C₁₋₁₂ alkyl, especially C₃₋₁₀ alkyl, C₂₋₆ alkenyl, C₂₋₆ alkynyl, C₃₋₈ cycloalkyl, C₁₋₆ alkyl, C₃₋₈ cycloalkyl, phenyl, phenyl-C₁₋₆ alkyl, especially benzyl, halophenyl-C₁₋₆ alkyl or pyridyl-C₁₋₆ alkyl group, or R¹ and R² together represent a saturated carbon chain containing three to eight carbon atoms while optionally one or more additional oxygen atoms may be present in the chain and which chain may optionally be aryl- or cycloalkyl-fused. Preferably R¹ and R² each independently represent a hydrogen atom, a C₂₋₁₂ alkyl, C₂₋₅ alkenyl, C₅₋₇ cyclo-C₁₋₂ alkyl, C₃₋₇ cycloalkyl or phenyl-C₁₋₂ alkyl group, or R₂ and R₃ together represent a saturated chain containing four or five carbon atoms while optionally additional oxygen atoms may be present and which chain optionally may be aryl- or cycloalkyl-fused, especially cyclopentyl, cyclohexyl or cycloheptyl fused, each of the above groups optionally substituted by one or more halogen atoms, especially chlorine and/or fluorine atoms, or C₁₋₄ alkyl, C₁₋₄ haloalkyl, C₄₋₆ cycloalkenyl or C₁₋₄ alkoxy groups.

[0023] In a preferred embodiment one of the radicals R¹ and R² is a branched C₃₋₁₂ alkyl or C₃₋₅ alkenyl group and the other is hydrogen or a straight chained C₁₋₄ alkyl group.

[0024] The invention also especially relates to compounds of formula I in which R² represents a hydrogen atom or a methyl group, preferably a hydrogen atom.

[0025] Another particular sub-group is that in which R¹ and R² each independently represent a hydrogen atom or a linear or branched C₁₋₁₂ alkyl group, especially a C₁₋₁₀ alkyl group, an allyl, C₃₋₇ cycloalkyl optionally fused with a cyclohexyl group, benzyl or phenyl group, or R¹ and R² together represent a saturated C₄₋₇ carbon chain, especially a C₄₋₆ carbon chain, which optionally may contain an additional oxygen atom and which optionally may be fused with a cyclohexyl ring, each of the above groups optionally substituted by a fluorine, chlorine or bromine atom or one or two methyl groups, a t-butyl, cyclohexyl, cyclohexenyl, phenyl or pyridyl group.

[0026] Particularly preferred are the compounds of the subgeneric formula IA,



wherein

R¹ and R² have the meaning given and

Hal represents halogen.

[0027] The compounds according to formula I are oils, gums, or, predominantly crystalline solid materials. They are superior through their valuable fungicidal properties. For example, they can be used in agriculture or related fields for the control of phytopathogenic fungi such as *Alternaria solani*, *Botrytis cinerea*, *Cercospora beticola*, *Cladosporium herbarum*, *Corticium rolfsii*, *Erysiphe graminis*, *Helminthosporium tritici repens*, *Leptosphaeria nodorum*, *Microneotriella nivalis*, *Monilinia fructigena*, *Mycosphaerella ligulicola*, *Mycosphaerella pinodes*, *Pycularia grisea f.sp. oryzae*, *Rhizoctonia solani* and *Sclerotinia sclerotiorum*, in particular for the control of *Pycularia grisea f.sp. oryzae* and *Rhizoctonia solani*. The compounds of formula I according to the invention possess a high fungicidal activity within a wide concentration range and may be used in agriculture without any difficulties.

[0028] Preferred compounds are those in which R² represents a bromine or chlorine atom.

[0029] Good results in terms of control of phytopathogenic fungi are obtained with a compound as defined in formula I wherein

R¹ represents straight chained or branched C₁₋₆-alkyl or straight chained or branched C₂₋₆-alkenyl, in particular branched C₃₋₆ alkyl or branched C₃₋₆ alkenyl, and

R² represents hydrogen or C₁-C₆-alkyl, in particular straight chained C₁-C₆-alkyl, or wherein

R¹ and R² together with the adjacent nitrogen atom represent a heterocyclic ring with 5 or 6 carbon atoms being optionally substituted with one or two C₁-C₆-alkyl groups, in particular a heterocyclic ring being optionally substituted by one or two methyl groups selected from pyrrolidiny, pyrrolidin, piperidiny, piperaziny and morpholin-4-yl.

[0030] Especially good results in terms of control of phytopathogenic fungi are obtained by using, for example, the following compounds of formula I:

5-chloro-6-(pentafluorophenyl)-7-(4-methyl-piperidin-1-yl)-[1,2,4]triazolo[1,5-a]pyrimidine;

5-chloro-6-(pentafluorophenyl)-7-(N-isopropylamino)-[1,2,4]triazolo[1,5-a]pyrimidine; and

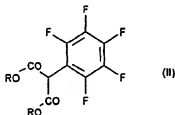
5-chloro-6-(pentafluorophenyl)-7-(N-ethyl-N-2-methylallyl-amino)-[1,2,4]triazolo[1,5-a]pyrimidine.

[0031] The conventional methods for preparing dialkylalkyl phenylmalonates, i.e. the reaction of alkyl phenyl-acetates with carboxylates or carbondioxide in the presence of strong bases, are not applicable for the preparation of dialkyl pentafluorophenylmalonates, since the strong bases will, at least in part, substitute the fluoro atoms of the pentafluorophenyl moiety.

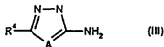
[0032] Therefore, the present invention further provides a process for the preparation of a compound of formula I as defined above which comprises

(a) reacting hexafluorobenzene with a dialkylmalonate in the presence of a base,

(b) treating the resulting 2-pentafluorophenylmalonate of formula II,

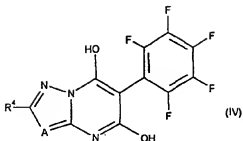


in which R is alkyl, aryl or benzyl, in particular C₁-C₆-alkyl, with a compound of formula III,



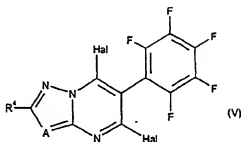
in which R⁴ and A have the meaning given in the preceding claims, in the presence of a base,

(c) treating the resulting dihydroxyazoloxy pyrimidine of formula IV,



in which R⁴ and A have the meaning given in the preceding claims, with a halogenating agent, and

(d) treating the resulting compound of the general formula V



in which

R⁴ and A are as defined in any one of the preceding claims; and
Hal represents a chlorine or bromine atom, with an amine of the general formula VI



in which

R¹ and R² are as defined in any one of the preceding claims, to produce a compound of formula I in which R³ represents a chlorine or bromine atom;

(e) if desired, reacting the compound of formula I formed in (d) with a fluorinating agent to produce a compound of formula I in which R³ represents a fluorine atom, and

(f) if desired, reacting the compound of formula I formed in (d) with ammonia and, subsequently, with diiodomethane in the presence of a diazotising agent to produce a compound of formula I in which R³ represents an iodine atom,

(g) if desired, reacting the compound of formula I formed in (d) with an amine of formula HNR⁵R⁶ to produce a compound of formula I in which R³ represents -NR⁵R⁶.

[0033] The process of step (a) is conveniently carried out in the presence of a solvent. Suitable solvents include polar aprotic solvents as for example sulfolane, dimethylformamide or a mixture thereof. The reaction is suitably carried out at a temperature in the range from room temperature (about 15 °C) to the reflux temperature of the reaction mixture, the preferred reaction temperature being from 40 °C to the reflux temperature of the reaction mixture. It is also preferred that the reaction is carried out in the presence of a base. Suitable bases include tertiary amines, such as triethylamine, and inorganic bases, such as potassium carbonate or sodium carbonate.

[0034] Compounds of formula IV can be prepared by reacting 3-amino-1,2,4-triazole (A = N) or 3-amino-1,2-diazole (A = CR²) of formula III with 2-penta-fluorophenyl-malonic acid ester under alkaline conditions according preferably using high boiling tertiary amines as for example tri-n-butylamine.

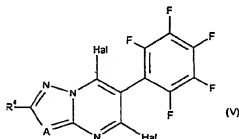
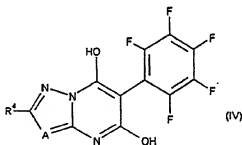
[0035] The process of step (c) is conveniently carried out with a brominating or chlorinating agent, such as phosphorus oxybromide or phosphorus oxychloride, neat or in the presence of a solvent. The reaction is suitably carried out at a temperature in the range from 0 °C to 150 °C, the preferred reaction temperature being from 80 °C to 125 °C.

[0036] The process of step (d) is conveniently carried out in the presence of a solvent. Suitable solvents include ethers, such as dioxane, diethyl ether and, especially, tetrahydrofuran, halogenated hydrocarbons such as dichloromethane and aromatic hydrocarbons, for example toluene. The reaction is suitably carried out at a temperature in the range from 0 °C to 70 °C, the preferred reaction temperature being from 10 °C to 35 °C. It is also preferred that the reaction is carried out in the presence of a base. Suitable bases include tertiary amines, such as triethylamine, and inorganic bases, such as potassium carbonate or sodium carbonate. Alternatively, an excess of the compound of formula III may serve as a base.

[0037] The process of step (e) is conveniently carried out in the presence of a solvent. Suitable solvents include sulfolane, dimethylformamide or a mixture of acetonitrile and a crown ether. If sulfolane or dimethylformamide is used as solvent, it is advantageous to use toluene as a co-solvent to aid dehydration of the fluorinating agent. The reaction is suitably carried out at a temperature in the range from room temperature (about 15 °C) to the reflux temperature of the reaction mixture, the preferred reaction temperature being from 40 °C to the reflux temperature of the reaction mixture. Suitable fluorinating agents include alkali metal fluorides, especially potassium fluoride, and antimony fluoride.

[0038] The first process of step (f) is conveniently carried out in the presence of a solvent. Suitable solvents include ethers, such as dioxane, diethyl ether and tetrahydrofuran, halogenated hydrocarbons such as dichloromethane and aromatic hydrocarbons such as toluene. The reaction is suitably carried out at a temperature in the range from 20 °C to the reflux temperature of the reaction mixture, the preferred reaction temperature being from 40 °C to the reflux temperature of the reaction mixture. It is also preferred that the reaction is carried out in the presence of a base. Suitable bases include tertiary amines, such as triethylamine, and inorganic bases, such as potassium carbonate or sodium carbonate. Alternatively an excess of ammonia may serve as base. For the diazotisation process conveniently the isolated material of the first process is used. The diazotising agent may be any alkyl ester of nitrous acid, isopentyl nitrite being especially preferred. If an alkyl ester of nitrous acid is used; this may serve as a co-solvent with the diiodomethane. The reaction is suitably carried out at a temperature from 60 °C to 120 °C, the preferred reaction temperature being from 70 °C to 110 °C.

[0039] The present invention further provides novel intermediates of formulae IV and V for the preparation of a compound of formula I as defined above:



in which R, R¹, A and Hal have the meaning given above.

[0040] The compounds of formula I have been found to have fungicidal activity. Accordingly the invention further provides a fungicidal composition which comprises a carrier and, as active ingredient, at least one compound of formula I as defined above. A method of making such a composition is also provided which comprises bringing a compound of formula I as defined above into association with at least one carrier. Such a composition may contain a single compound or a mixture of several compounds of the present invention. It is also envisaged that different isomers or mixtures of isomers may have different levels or spectra of activity and thus compositions may comprise individual isomers or mixtures of isomers.

[0041] A composition according to the invention preferably contains from 0.5% to 95% by weight of active ingredient.

[0042] A carrier in a composition according to the invention is any material with which the active ingredient is formulated to facilitate application to the locus to be treated, which may for example be a plant, seed or soil, or to facilitate storage, transport or handling. A carrier may be a solid or a liquid, including material which is normally gaseous but which has been compressed to form a liquid, and any of the carriers normally used in formulating fungicidal compositions may be used.

[0043] The compositions may be manufactured into e.g. emulsion concentrates, solutions which may be sprayed directly or diluted, diluted emulsions, wettable powders, soluble powders, dusts, granulates, water-dispersible granulates, microencapsulates by well-established procedures. The form of application such as spraying, atomizing, dispersing, pouring may be chosen like the compositions according to the desired objectives and the given circumstances.

[0044] The formulations, i.e. the compositions which comprise at least one compound according to general formula I and optionally solid and/or liquid auxiliaries and adjuvants, may be prepared by well-established procedures, e.g. intensive mixing and/or grinding of the active ingredients with other substances, such as fillers, solvents, solid carriers, and optionally surface-active compounds (tensides).

[0045] Solvents may be aromatic hydrocarbons, preferably the fractions C₈ to C₁₂, e.g. xylenes or xylene mixtures, substituted naphthalenes, phthalic acid esters, such as dibutyl or dioctyl phthalate, aliphatic hydrocarbons, e.g. cyclohexane or paraffins, alcohols and glycols as well as their ethers and esters, e.g. ethanol, ethylene glycol mono- and dimethyl ether, ketones such as cyclohexanone, strongly polar solvents such as N-methyl 2-pyrrolidone, dimethyl sulfoxide, alkyl formamides, epoxidized vegetable oils, e.g. epoxidized coconut or soybean oil, water. Mixtures of different liquids are often suitable.

[0046] Solid carriers, which may be used for dusts or dispersible powders, may be mineral fillers, such as calcite, talc, kaolin, montmorillonite, attapulgite. The physical properties may be improved by addition of highly dispersed silica gel or highly dispersed polymers. Carriers for granulates may be porous material, e.g. pumice, broken brick, sepiolite, bentonite, non-sorptive carriers may be calcite or sand. Additionally, a multitude of pre-granulated inorganic or organic materials may be used, such as dolomite or crushed plant residues.

[0047] Fungicidal compositions are often formulated and transported in concentrated form which is subsequently diluted by the user before application. The presence of small amounts of a carrier which is a surface-active agent facilitates this process of dilution. Thus, preferably one carrier in a composition according to the invention is a surface active agent. For example, the composition may contain at least two carriers, at least one of which is a surface active agent.

[0048] Suitable surface-active substances may be non-ionic, anionic or cationic tensides with good dispersing, emulating and wetting properties depending on the nature of the compound according to general formula I to be formulated. Tensides may also mean mixtures of tensides.

[0049] Suitable tensides may be so-called water-soluble soaps as well as water-soluble synthetic surface-active compounds.

[0050] Soaps usually are alkali, earth alkali or optionally-substituted ammonium salts of higher fatty acids (C₁₀-C₂₀), e.g. the sodium or potassium salts of oleic or stearic acid or of mixtures of natural fatty acids which are prepared, for example, from coconut or tallow oil. Furthermore, methyl-taurine salts of fatty acids may be used.

[0051] However, so-called synthetic tensides are preferably used, especially fatty sulphonates, fatty sulphates, sulphonated benzimidazole derivatives or alkyl aryl sulphonates.

[0052] The fatty sulphates or fatty sulphonates are normally used as alkali, earth alkali or optionally-substituted ammonium salts and have an alkyl moiety of 8 to 22 carbon atoms, whereby alkyl also means the alkyl moiety of acyl residues, such as the sodium or calcium salt of lignin sulphonic acid, of sulphuric acid dodecylate or of a mixture of fatty alcohols prepared from natural fatty acids. This also includes the salts of sulphuric acid esters, sulphonic acids and adducts of fatty alcohols and ethylene oxide. The sulphonated benzimidazole derivatives preferably contain 2 sulphonic acid residues and a fatty acid residue with 8 to 22 carbon atoms. Alkyl aryl sulphonates are, for example, the sodium, calcium or triethyl ammonium salts of dodecyl benzene sulphonic acid, dibutyl naphthalene sulphonic acid or of a condensate of naphthalene sulphonic acid and formaldehyde.

[0053] Furthermore, phosphates, such as the salts of the phosphoric acid ester of *ap*-nonylphenol-(4-14)-ethylene oxide adduct or phospholipids, may be used.

[0054] Non-ionic tensides are preferably polyglycoether derivatives of aliphatic or cycloaliphatic alcohols, saturated or non-saturated fatty acids and alkylphenols, which have 3 to 10 glycol ether groups and 8 to 20 carbon atoms in the (aliphatic) hydrocarbon residue and 6 to 18 carbon atoms in the alkyl residue of the alkyl phenols.

[0055] Other suitable non-ionic tensides are the water-soluble, 20 to 250 ethylene glycol ether groups containing polyadducts of ethylene oxide and polypropylene glycol, ethylene diamino polypropylene glycol and alkyl polypropylene glycol with 1 to 10 carbon atoms in the alkyl moiety, the substances normally contain 1 to 5 ethylene glycol units per propylene glycol unit.

[0056] Examples of non-ionic tensides are nonylphenol polyethoxy ethanol, castor oil polyglycol ether, polyadducts of ethylene oxide and polypropylene, tributyl phenoxy polyethoxy ethanol, polyethylene glycol, octyl phenoxy polyethoxy ethanol.

[0057] Furthermore, fatty acid esters of polyoxy ethylene sorbitan, such as polyoxy ethylene sorbitan trioleate may be used.

[0058] Cationic tensides preferably are quaternary ammonium salts, which have at least one alkyl residue with 8 to 22 carbon atoms and, furthermore, low, optionally-halogenated alkyl, benzyl or hydroxyalkyl residues. The salts are preferably halides, methyl sulphates or alkyl sulphates, e.g. stearyl trimethyl ammonium chloride or benzyl bis(2-chloroethyl) ethyl ammonium bromide.

[0059] The compositions of the invention may for example be formulated as wettable powders, dusts, granules, solutions, emulsifiable concentrates emulsions, suspension concentrates and aerosols. Wettable powders usually contain 25%, 50% or 75% w/w of active ingredient and usually contain in addition to solid inert carrier, 3%-10% w/w of a dispersing agent and, where necessary, 0%-10% w/w of stabiliser(s) and/or other additives such as penetrants or stickers. Dusts are usually formulated as a dust concentrate having a similar composition to that of a wettable powder but without a dispersant, and may be diluted in the field with further solid carrier to give a composition usually containing 0.5%-10% w/w of active ingredient. Granules are usually prepared to have a size between 10 and 100 mesh ASTM (approx. 2.00 mm - 0.15 mm), and may be manufactured by agglomeration or impregnation techniques. Generally, granules will contain 0.5%-75% active ingredient and 0-10% w/w of additives such as stabiliser, surfactants, slow release modifiers and binding agents. The so called "dry flowable powders" consist of relatively small granules having a relatively high concentration of active ingredient. Emulsifiable concentrates usually contain, in addition to a solvent or a mixture of solvents, 1%-50% w/v active ingredient, 2%-20% w/v emulsifiers and 0%-20% w/v of other additives such as stabilisers, penetrants and corrosion inhibitors. Suspension concentrates are usually compounded so as to obtain a stable, non-sedimenting flowable product and usually contain 10%-75% w/w active ingredient, 0.5%-15% w/w of dispersing agents, 0.1%-10% w/w of suspending agents such as protective colloids and thixotropic agents, 0%-10% of other additives such as defoamers, corrosion inhibitors, stabilisers, penetrants and stickers, and water or an organic liquid in which the active ingredient is substantially insoluble; certain organic solids or inorganic salts may be present. dissolved in the formulation to assist in preventing sedimentation or as antifreeze agents for water.

[0060] Aqueous dispersions and emulsions, for example compositions obtained by diluting a wettable powder or a concentrate according to the invention with water, also lie within the scope of the invention. The said emulsions may be of the water-in-oil or of the oil-in-water type, and may have a thick 'mayonnaise' like consistency.

[0061] The composition of the invention may also contain other ingredients, for example other compounds possessing herbicidal, insecticidal or fungicidal properties.

[0062] Of particular interest in enhancing the duration of the protective activity of the compounds of this invention is the use of a carrier which will provide slow release of the fungicidal compounds into the environment of a plant which is to be protected. Such slow-release formulations could, for example, be inserted in the soil adjacent to the roots of a plant, or could include an adhesive component enabling them to be applied directly to the stem of a plant.

[0063] As commodity the compositions may preferably be in a concentrated form whereas the end-user generally employs diluted compositions. The compositions may be diluted to a concentration of 0.001% of active ingredient (a.i.). The doses usually are in the range from 0.01 to 10 kg a.i./ha.

[0064] The invention still further provides the use as a fungicide of a compound of formula I as defined above or a composition as defined above, and a method for combating fungus at a locus, which comprises treating the locus, which may be for example plants subject to or subjected to fungal attack, seeds of such plants or the medium in which such plants are growing or are to be grown, with such a compound or composition.

[0065] The present invention is of wide applicability in the protection of crop and ornamental plants against fungal attack. Typical crops which may be protected include vines, grain crops such as wheat and barley, rice, sugar beet, top fruit, peanuts, potatoes and tomatoes. The duration of the protection is normally dependent on the individual compound selected, and also a variety of external factors, such as climate, whose impact is normally mitigated by the use of a suitable formulation.

[0066] The following examples further illustrate the present invention.

Example 1**(1a)** Diethyl pentafluorophenyl-malonate

- 5 **[0067]** A mixture of N,N-dimethylformamide (35 mL) and potassium carbonate (7.52 g, 54 mmol) is heated to 150 °C and diethyl malonate (6.45 g, 53 mmol) is added without further heating. Hexafluorobenzene (10 g, 53 mmol) is added to the reaction mixture at 115 °C. The resulting reaction mixture is stirred for 3 hours at 120 °C. The hot solution is poured onto crushed ice (160 mL), carefully acidified with sulfuric acid and cooled to 0 °C. The aqueous layer is removed and the residue is dissolved in diethyl ether. The ethereal solution is washed with aqueous sodium hydrogencarbonate and water and dried with sodium sulfate. The ether is distilled off *in vacuo*. The resulting product (9.5 g) contains 75 % diethyl pentafluorophenyl malonate and is used as intermediate without further purification.
- 10 ¹H-NMR data (CDCl₃/tetramethylsilane):
s (5.4 ppm), m (4.3-4.1 ppm), m (1.3-1.1 ppm).

- 15 **(1b)** 5,7-Dihydroxy-6-pentafluorophenyl-1,2,4-triazolo[1.5a]-pyrimidine

- [0068]** A mixture of (1a) (22 mmol), tributylamine (5.71 mL, 24 mmol) and 2-amino-1,2,4-triazole (1.65 g, 22 mmol) is heated to 180 °C for 6 hours. The reaction mixture is cooled down to 50 °C and an aqueous solution of sodium hydroxide (2.2 g in 25 mL) is added. The mixture is stirred for 30 minutes, the aqueous layer is separated, extracted with ether twice and acidified with concentrated hydrochloric acid. The resulting white solid is filtered off, washed with water and diisopropyl ether and dried. The resulting yellowish powder (5.2 g) decomposes at 200 °C.

25

Elemental analysis:			
	% C	% H	% N
calculated	41.48	0.94	17.59
found	39.52	1.62	17.53

- 30 **(1c)** 5,7-Dichloro-6-pentafluorophenyl-1,2,4-triazolo[1.5a]-pyrimidine

- [0069]** A mixture of (1b) (5.15 g, 16 mmol) and phosphorousoxychloride (20 mL) is refluxed for 4 hours at 120 °C. Dichloromethane (100 mL) is added to the reaction mixture upon cooling to room temperature. After adding water (125 mL) at temperatures below 40 °C, the organic phase is separated, dried with sodium sulfate and the solvent is distilled off. The resulting oil (2.6 g) is used as intermediate without further purification.

- 35 ¹H-NMR data (CDCl₃ / tetramethylsilane):
s (8.95 ppm).

(1d) 5-Chloro-7-N-isopropylamino-6-pentafluorophenyl-1,2,4-triazolo[1.5a]pyrimidine.

- 40 **[0070]** A mixture of isopropylamine (1.4 mmol), triethylamine (1.4 mmol) and dichloromethane (10 mL) is added to a mixture of (1c) (1.4 mmol) and dichloromethane (30 mL) under stirring. The reaction mixture is stirred 18 hours at room temperature, subsequently washed two times with 1 N hydrochloric acid and once with water. The organic layer is separated, dried with anhydrous sodium sulphate and the solvent is evaporated under reduced pressure.
- 45 Treatment of the resulting light brown oil with *tert*-butyl methyl ether (50 mL) yields white crystals (79% of th.) having a melting point of 167-168 °C.

Examples 2-33

- 50 **[0071]** The following examples (Table I; structure and melting point) are synthesized analogously to Example 1.

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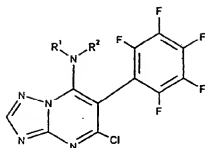


TABLE I

Example	R ¹	R ²	mp (°C)
2	-(CH ₂) ₂ -CHCH ₃ -(CH ₂) ₂ -		173-174
3	2-methylallyl	ethyl	100-102
4	cyclopentyl	H	145-146
5	ethyl	ethyl	118-119
6	allyl	ethyl	156
7	H	H	
8	cyclohexyl	H	178-179
9	3,3,4-trimethylcyclopentyl	H	
10	-(CH ₂) ₂ -CHCl-(CH ₂) ₂ -		
11	methyl	H	
12	ethyl	H	
13	methyl	methyl	
14	-(CH ₂) ₆ -		
15	-(CH ₂) ₅ -		164
16	-(CH ₂) ₄ -		
17	-CH ₂ -CHCH ₃ -(CH ₂) ₃ -		146
18	-CHCH ₃ -(CH ₂) ₄ -		
19	-(CH ₂) ₂ -O-(CH ₂) ₂ -		
20	1-ethylpropyl	H	
21	-CH ₂ -CHCH ₃ -CH ₂ -CHCH ₃ -CH ₂ -		
22	-CH ₂ -CH=CH-(CH ₂) ₂ -		
23	-(CH ₂) ₂ -N(CH ₃)-(CH ₂) ₂ -		
24	-CH ₂ -C(CH ₃) ₂ -(CH ₂) ₃ -		
25	-CHCH ₃ -(CH ₂) ₃ -		
26	cyclopropyl	H	214
27	norbornyl	H	194-195
28	1-methylpropyl	H	122
29	2-methylpropyl	H	169
30	cycloheptyl	H	155
31	allyl	methyl	140-141
32	2-methylpropyl	methyl	122
33	1,1,3,3-tetramethylbutyl	H	

Biological Investigations

Determination of Minimum Inhibitory concentration by Test Compounds in the Serial Dilution Test with Various Phytopathogenic Fungi

[0072] The MIC (Minimum Inhibitory Concentration) value, which indicates the lowest concentration of the active ingredient in the growth medium which causes a total inhibition of mycelial growth, is determined by serial dilution tests using Microtiter plates with 24 or 48 wells per plate. The dilution of the test compounds in the nutrient solution and the distribution to the wells is carried out by a TECAN RSP 5000 Robotic Sample Processor. The following test compound concentrations are used: 0.05, 0.10, 0.20, 0.39, 0.78, 1.56, 3.13, 6.25, 12.50, 25.00, 50.00 and 100.00 mg/mL. For preparation of the nutrient solution, V8 vegetable juice (333 mL) is mixed with calcium carbonate (4.95 g), centrifuged, the supernatant (200 mL) diluted with water (800 mL) and autoclaved at 121 °C for 30 min.

[0073] The respective inocula (*Alternaria solani*, ALTESO; *Botrytis cinerea*, BOTRICI; *Leptosphaeria nodorum*, LEP-TNO; *Phytophthora infestans*, PHYTIN; *Magnaporthe grisea oryzae*, PYRIOR; *Pyrenophora teres*, PYRANTE; *Rhizoctonia solani*, RHIZSO;) are added into the wells as spore suspensions (50 mL; 5x10⁹/mL) or agar slices (6 mm) of an agar culture of the fungus.

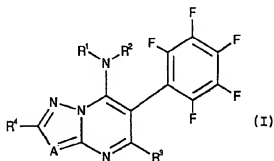
[0074] After 6-12 days incubation at suitable temperatures (18-25°C), the MIC values are determined by visual inspection of the plates (Table II; n.l. = not tested).

Table II

Ex. No.	ALTESO	BOTRICI	LEPTNO	PHYTIN	PYRIOR	PYRANTE	RHIZSO
1	12.5	12.5	25	100	1.56	25	6.25
2	0.2	0.78	6.25	110	0.1	1.56	0.78
3	1.56	0.78	3.13	110	<0.05	1.56	0.39
4	0.4	6.25	50	110	3.13	3.13	110
5	12.5	3.13	50	100	0.78	3.13	25
6	6.25	1.56	50	50	0.78	3.13	0.78
8	3.13	6.25	110	110	25	0.78	110
15	0.4	1.56	12.5	100	0.78	1.56	110
17	3.13	3.13	110	110	3.13	1.56	110
27	0.4	3.13	110	100	25	0.4	110
28	6.25	12.5	110	100	3.13	12.5	6.25
31	1.56	1.56	100	110	0.78	1.56	6.25
32	6.25	3.13	110	100	0.78	1.56	1.56

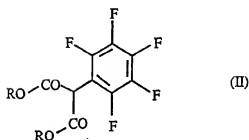
Claims

1. A compound of formula I

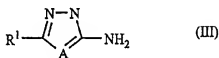


in which

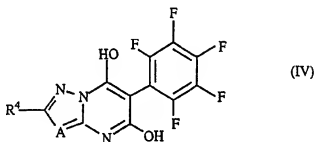
- 5 R^1 and R^2 each independently represent hydrogen or C_1 - C_{10} -alkyl, C_2 - C_{10} -alkenyl, C_3 - C_{10} -alkynyl, C_3 - C_{10} -alkadienyl, C_6 -, C_{10} - or C_{14} -aryl, heteroaryl having 5 or 6 ring atoms selected from carbon, nitrogen, oxygen and sulphur, at least one of which being nitrogen, oxygen or sulphur, C_3 - C_6 -cycloalkyl, saturated heterocyclyl having 5 or 6 ring atoms selected from carbon, nitrogen, oxygen and sulphur, at least one of which being nitrogen, oxygen or sulphur, or
- 10 R^1 and R^2 together with the adjacent nitrogen atom represent a heterocyclic ring, wherein each group is optionally substituted by one or more halogen atoms or nitro, cyano, C_{1-6} -alkyl, C_{3-6} -cycloalkyl, C_{3-6} -cycloalkenyl, C_{1-6} -haloalkyl, C_{3-6} -halocycloalkyl, C_{1-6} -alkoxy, C_{1-6} -haloalkoxy, phenyl, halo- or dihalo-phenyl or pyridyl.
- 15 R^3 represents a halogen atom or a group $-NR^5R^6$, wherein R^5 and R^6 each independently have one of the meanings given for R^1 and R^2 .
- R^4 represents hydrogen or C_1 - C_{10} -alkyl or C_6 -, C_{10} - or C_{14} -aryl; and
- A represents N or CR^7 , wherein R^7 has the meaning given for R^4 .
- 20 2. A compound according to Claim 1 in which
- R^3 represents a bromine or chlorine atom.
- 25 3. A compound according to Claim 1 or 2 in which R^1 represents straight chained or branched C_1 - C_6 -alkyl or straight chained or branched C_2 - C_6 -alkenyl, and R^2 represents hydrogen or C_1 - C_6 -alkyl, or R^1 and R^2 together with the adjacent nitrogen atom represent a heterocyclic ring with 5 or 6 carbon atoms being optionally substituted with one or two C_1 - C_6 -alkyl groups.
- 30 4. A compound according to any one of the preceding claims in which A is N, and R^4 is hydrogen.
5. A compound according to any one of the preceding claims in which R^1 and R^2 together with the adjacent nitrogen atom represent heterocyclic group selected from 4-methyl-piperidin-1-yl, 2-methyl-piperidin-1-yl, 3,6-dihydro-2H-pyridin-1-yl, 2-ethyl-piperidin-1-yl and azepan-1-yl.
- 35 6. A compound according to any one of the preceding claims, wherein the compound is selected from the group consisting of:
- 5-chloro-6-(pentafluorophenyl)-7-(4-methyl-piperidin-1-yl)-[1,2,4]triazolo[1,5-a]pyrimidine
 5-chloro-6-(pentafluorophenyl)-7-(N-isopropylamino)-[1,2,4]triazolo[1,5-a]pyrimidine
 5-chloro-6-(pentafluorophenyl)-7-(N-ethyl,N-2-methylallylamino)-[1,2,4]triazolo[1,5-a]pyrimidine
- 40 7. A process for the preparation of a compound of formula I as defined in any one of the preceding claims which comprises
- 45 (a) reacting hexafluorobenzene with a dialkylmalonate in the presence of a base,
- (b) treating the resulting 2-pentafluorophenylmalonate of formula II.
- 50
- 55



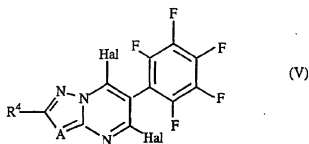
15 in which R is alkyl, aryl or benzyl, with a compound of formula III,



25 in which R⁴ and A have the meaning given in any one of the preceding claims, in the presence of a base,
(c) treating the resulting dihydroxyazolopyrimidine of formula IV,

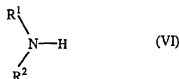


40 in which R⁴ and A have the meaning given in any one of the preceding claims, with a halogenating agent, and
(d) treating the resulting compound of the general formula V



in which
R⁴ and A are as defined in any one of the preceding claims; and

Hal represents a chlorine or bromine atom, with an amine of the general formula VI



in which

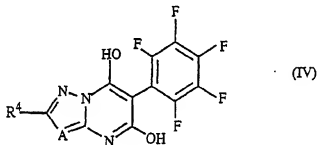
R^1 and R^2 are as defined in any one of the preceding claims, to produce a compound of formula I in which R^3 represents a chlorine or bromine atom; and

(e) if desired, reacting the compound of formula I formed in (d) with a fluorinating agent to produce a compound of formula I in which R^3 represents a fluorine atom; or

(f) if desired, reacting the compound of formula I formed in (d) with ammonia and, subsequently, with diiodomethane in the presence of a diazotising agent to produce a compound of formula I in which R^3 represents an iodine atom; or

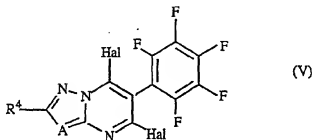
(g) if desired, reacting the compound of formula I formed in (d) with an amine of formula NHR^5R^6 to produce a compound of formula I in which R^3 represents $-NR^5R^6$.

8. A compound of formula IV



in which R^4 and A have the meaning given in any one of the preceding claims.

9. A compound of formula V



in which R^4 and A are as defined in any one of the preceding claims; and Hal represents a chlorine or bromine atom.

10. A fungicidal composition which comprises a carrier, and as active agent, at least one compound of formula I as

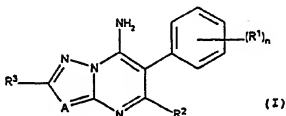
defined in any one of Claims 1 to 6.

11. A method of combating fungus at a locus which comprises treating the locus with a compound of formula I as defined in any one of Claims 1 to 6 or with a composition as defined in Claim 10.

12. The use as a fungicide of a compound of formula I as defined in any one of Claims 1 to 6 or a composition as defined in Claim 10.

Patentansprüche

1. Verbindungen der Formel I



in welcher

R^1 und R^2 jeweils unabhängig voneinander für Wasserstoff oder C_1 - C_{10} -Alkyl, C_2 - C_{10} Alkenyl, C_2 - C_{10} -Alkynyl, C_4 - C_{10} -Alkadienyl, C_6 -, C_{10} - oder C_{14} -Aryl, Heteroaryl mit 5 oder 6 Ringatomen ausgewählt aus Kohlenstoff, Stickstoff, Sauerstoff und Schwefel, von denen wenigstens eines Stickstoff, Sauerstoff oder Schwefel ist, C_3 - C_6 -Cycloalkyl, gesättigtes Heterocyclyl mit 5 oder 6 Ringatomen ausgewählt aus Kohlenstoff, Stickstoff, Sauerstoff und Schwefel, von denen wenigstens eines Stickstoff, Sauerstoff oder Schwefel ist, stehen, oder

R^1 und R^2 zusammen mit dem benachbarten Stickstoffatom für einen heterocyclischen Ring stehen, wobei jede Gruppe gegebenenfalls durch ein oder mehrere Halogenatome oder Nitro, Cyano, C_{1-5} -Alkyl, C_{3-6} -Cycloalkyl, C_{3-6} -Cycloalkenyl, C_{1-6} -Halogenalkyl, C_{3-6} -Halogenocycloalkyl, C_{1-5} -Alkoxy, C_{1-5} -Halogenalkoxy, Phenyl, Halogen- oder Dihalogenphenyl oder Pyridyl substituiert ist,

R^3 für ein Halogenatom oder eine Gruppe $-NR^5R^6$ steht, wobei R^5 und R^6 jeweils unabhängig voneinander eine der für R^1 und R^2 angegebenen Bedeutungen haben,

R^4 für Wasserstoff oder C_1 - C_{10} -Alkyl oder C_6 -, C_{10} oder C_{14} -Aryl steht, und

A für N oder CR^7 steht, wobei R^7 die für R^4 angegebene Bedeutung hat.

2. Verbindungen nach Anspruch 1, in welchen

R^3 für ein Brom- oder Chloratom steht.

3. Verbindungen nach Anspruch 1 oder 2, in welchen R^1 für geradkettiges oder verzweigtes C_1 - C_6 -Alkyl oder geradkettiges oder verzweigtes C_2 - C_6 -Alkenyl steht und R^2 für Wasserstoff oder C_1 - C_6 -Alkyl steht, oder R^1 und R^2 zusammen mit dem benachbarten Stickstoffatom für einen heterocyclischen Ring mit 5 oder 6 Kohlenstoffatomen stehen, der gegebenenfalls durch eine oder zwei C_1 - C_6 -Alkylgruppen substituiert ist.

4. Verbindungen nach einem der vorhergehenden Ansprüche, in welchen A für N und R^4 für Wasserstoff steht.

5. Verbindungen nach einem der vorhergehenden Ansprüche, in welchen R^1 und R^2 zusammen mit dem benachbarten Stickstoffatom für eine aus 4-Methylpiperidin-1-yl, 2-Methylpiperidin-1-yl, 3,6-Dihydro-2H-pyridin-1-yl,

2-Ethylpiperidin-1-yl und Azepan-1-yl ausgewählte heterocyclische Gruppe stehen.

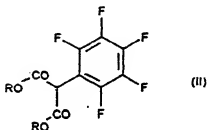
6. Verbindungen nach einem der vorhergehenden Ansprüche, ausgewählt aus der folgenden Gruppe:

- 5 5-Chlor-6-(pentafluorphenyl)-7-(4-methylpiperidin-1-yl)-[1,2,4]triazolo[1,5-a]pyrimidin
 5-Chlor-6-(pentafluorphenyl)-7-(N-isopropylamino)-[1,2,4]triazolo[1,5-a]pyrimidin
 5-Chlor-6-(pentafluorphenyl)-7-(N-ethyl,N-2-methylallylamino)-[1,2,4]triazolo[1,5-a]pyrimidin.

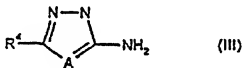
7. Verfahren zur Herstellung von den in einem der vorhergehenden Ansprüche definierten Verbindungen der Formel I, bei dem man

(a) Hexafluorbenzol in Gegenwart einer Base mit Malonsäuredialkylester umsetzt,

(b) das so erhaltene 2-Pentafluorphenylmalonat der Formel II

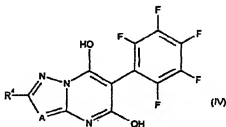


in welcher R für Alkyl, Aryl oder Benzyl steht, in Gegenwart einer Base mit einer Verbindung der Formel III



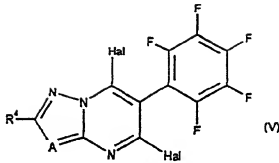
in welcher R4 und A die in einem der vorhergehenden Ansprüche angegebene Bedeutung haben, behandelt,

(c) das so erhaltene Dihydroxyazoloxyrimidin der Formel IV

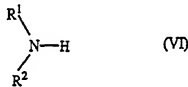


in welcher R4 und A die in einem der vorhergehenden Ansprüche angegebene Bedeutung haben, mit einem Halogenierungsmittel behandelt, und

(d) die so erhaltene Verbindung der allgemeinen Formel V



15 in welcher
 R⁴ und A wie in einem der vorhergehenden Ansprüche definiert sind,
 und
 Hal für ein Chlor- oder Bromatom steht, mit einem Amin der allgemeinen Formel VI



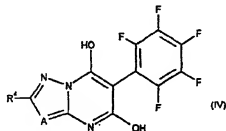
30 in welcher
 R¹ und R² wie in einem der vorhergehenden Ansprüche definiert sind,
 behandelt und so eine Verbindung der Formel I erhält, in welcher R³ für ein Chlor- oder Bromatom steht; und

35 (e) gewünschtenfalls die in (d) gebildete Verbindung der Formel I mit einem Fluorierungsmittel zu einer Verbindung der Formel I umsetzt, in der R³ für ein Fluoratom steht; oder

40 (f) gewünschtenfalls die in (d) gebildete Verbindung der Formel I mit Ammoniak und anschließend mit Diodmethan in Gegenwart eines Diazotierungsmittels zu einer Verbindung der Formel I umsetzt, in der R³ für ein Iodatom steht; oder

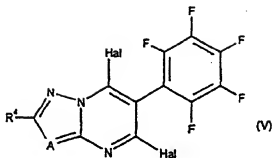
(g) gewünschtenfalls die in (d) gebildete Verbindung der Formel I mit einem Amin der Formel NHR⁵R⁶ zu einer Verbindung der Formel I umsetzt, in der R³ für -NR⁵R⁶ steht.

8. Verbindungen der Formel IV



in welcher R⁴ und A die in einem der vorhergehenden Ansprüche angegebene Bedeutung haben.

9. Verbindungen der Formel V



in welcher R^4 und A wie in einem der vorhergehenden Ansprüche definiert sind, und Hal für ein Chlor- oder Bromatom steht.

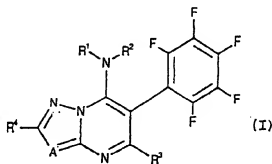
10. Fungizide Zusammensetzung, enthaltend einen Trägerstoff und, als Wirkstoff, wenigstens eine wie in einem der Ansprüche 1 bis 6 definierte Verbindung der Formel I.

11. Verfahren zur Bekämpfung von Pilzen an einem Ort, bei dem man den Ort mit einer wie in einem der Ansprüche 1 bis 6 definierten Verbindung der Formel I oder mit einer wie in Anspruch 10 definierten Zusammensetzung behandelt.

12. Verwendung einer wie in einem der Ansprüche 1 bis 6 definierten Verbindung der Formel I oder einer wie in Anspruch 10 definierten Zusammensetzung als Fungizid.

Revendications

1. Composé de formule I



dans laquelle

R^1 et R^2 représentent chacun indépendamment l'hydrogène ou un groupe alkyle en C_1-C_{10} , alcényle en C_2-C_{10} , alcylnyle en C_2-C_{10} , alcadiényle en C_4-C_{10} , aryle en C_6 , C_{10} ou C_{14} , hétéroaryle ayant 5 ou 6 atomes cycliques choisis parmi le carbone, l'azote, l'oxygène et le soufre, dont l'un au moins est un atome d'azote, d'oxygène ou de soufre, cycloalkyle en C_3-C_6 , hétérocyclyle saturé ayant 5 ou 6 atomes cycliques choisis parmi le carbone, l'azote, l'oxygène et le soufre, dont l'un au moins est un atome d'azote, d'oxygène ou de soufre, ou bien

R^1 et R^2 , avec l'atome d'azote adjacent, représentent un hétérocycle, chaque groupe étant facultativement substitué par un ou plusieurs atomes d'halogène ou groupes nitro, cyano, alkyle en C_1-C_6 , cycloalkyle

en C₃-C₆, cycloalcényle en C₃-C₆, halogénoalkyle en C₁-C₆, halogénocycloalkyle en C₃-C₆, alcoxy en C₁-C₆, halogénoalcoxy en C₁-C₆, phényle, halogéno- ou dihalogéno-phényle ou pyridyle, représente un atome d'halogène ou un groupe -NR⁵R⁶, où R⁵ et R⁶ ont chacun indépendamment

5 R⁴ représente l'hydrogène ou un groupe alkyle en C₁-C₁₀ ou aryle en C₆, C₁₀ ou C₁₄; et A représente N ou CR⁷, où R⁷ a la signification donnée pour R⁴.

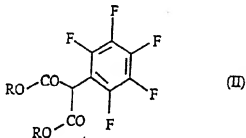
2. Composé selon la revendication 1, dans lequel R³ représente un atome de brome ou de chlore.
- 10 3. Composé selon la revendication 1 ou 2, dans lequel R¹ représente un groupe alkyle en C₁-C₆ à chaîne droite ou ramifiée ou alcényle en C₂-C₆ à chaîne droite ou ramifiée, et R² représente l'hydrogène ou un groupe alkyle en C₁-C₆, ou bien R¹ et R², avec l'atome d'azote adjacent, représentent un hétérocycle ayant 5 ou 6 atomes de carbone facultativement substitué par un ou deux groupes alkyle en C₁-C₆.
- 15 4. Composé selon l'une quelconque des revendications précédentes, dans lequel A est N, et R⁴ est l'hydrogène.
5. Composé selon l'une quelconque des revendications précédentes, dans lequel R¹ et R², avec l'atome d'azote adjacent, représentent un groupe hétérocyclique choisi parmi les groupes 4-méthylpipéridine-1-yle, 2-méthylpipéridine-1-yle, 3,6-dihydro-2H-pyridine-1-yle, 2-éthylpipéridine-1-yle et azépane-1-yle.
- 20 6. Composé selon l'une quelconque des revendications précédentes, dans lequel le composé est choisi dans la classe formée par les suivants :

5-chloro-6-(pentafluorophényl)-7-(4-méthylpipéridine-1-yl)-[1,2,4]triazolo[1,5-a]pyrimidine

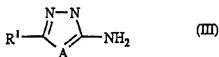
25 5-chloro-6-(pentafluorophényl)-7-(N-isopropylamino)-[1,2,4]triazolo[1,5-a]pyrimidine

5-chloro-6-(pentafluorophényl)-7-(N-éthyl-N-2-méthylallylamino)-[1,2,4]triazolo[1,5-a]pyrimidine.
7. Procédé pour la préparation d'un composé de formule I tel que défini dans l'une quelconque des revendications précédentes, qui consiste à

- (a) faire réagir de l'hexafluorobenzène avec un malonate de dialkyle en présence d'une base,
- (b) traiter le 2-pentafluorophénylmalonate résultant de formule II

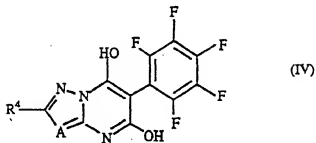


dans laquelle R est un groupe alkyle, aryle ou benzyle, avec un composé de formule III



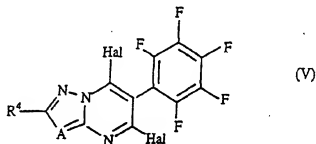
dans laquelle R⁴ et A sont tels que définis dans l'une quelconque des revendications précédentes, en présence d'une base,

(c) traiter la dihydroxyazolopyrimidine résultante de formule IV



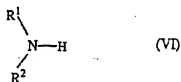
dans laquelle R^4 et A sont tels que définis dans l'une quelconque des revendications précédentes, avec un agent d'halogénéation, et

(d) traiter le composé résultant de formule générale V



dans laquelle

R^4 et A sont tels que définis dans l'une quelconque des revendications précédentes ; et
Hal représente un atome de chlore ou de brome,
avec une amine de formule générale VI



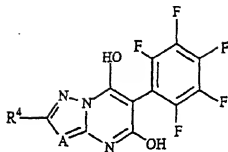
dans laquelle R^1 et R^2 sont tels que définis dans l'une quelconque des revendications précédentes,
pour produire un composé de formule I dans lequel R^3 représente un atome de chlore ou de brome ; et

(e) facultativement, faire réagir le composé de formule I formé en (d) avec un agent de fluoration pour produire un composé de formule I dans lequel R^3 représente un atome de fluor ; ou

(f) facultativement, faire réagir le composé de formule I formé en (d) avec l'ammoniac et, ensuite, avec le diiodométhane en présence d'un agent de diazotation pour produire un composé de formule I dans lequel R^3 représente un atome d'iode ; ou

(g) facultativement, faire réagir le composé de formule I formé en (d) avec une amine de formule $NH-R^5R^6$ pour produire un composé de formule I dans lequel R^3 représente $-NR^5R^6$.

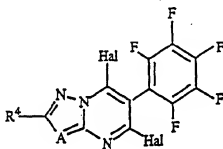
8. Composé de formule IV



(IV)

dans laquelle R⁴ et A sont tels que définis dans l'une quelconque des revendications précédentes.

9. Composé de formule V



(V)

dans laquelle

R⁴ et A sont tels que définis dans l'une quelconque des revendications précédentes ; et
Hal représente un atome de chlore ou de brome.

10. Composition fongicide qui comprend un support et, comme agent actif, au moins un composé de formule I tel que défini dans l'une quelconque des revendications 1 à 6.

11. Procédé pour combattre un champignon en un site, qui comprend le traitement du site avec un composé de formule I tel que défini dans l'une quelconque des revendications 1 à 6 ou avec une composition tel que définie dans la revendication 10.

12. Utilisation, comme fongicide, d'un composé de formule I tel que défini dans l'une quelconque des revendications 1 à 6 ou d'une composition telle que définie dans la revendication 10.